## Specification

Please amend the specification to recite the following replacement paragraphs. A marked-up version showing changes made to the specification through the replacement paragraphs is contained herewith as Appendix B.

## Please replace paragraph [0021] with the following replacement paragraph:

--[0021] FIGURES 1A-B depict the nucleotide sequence (SEQ ID NO:1), amino acid translation (SEQ ID NO:2) and intron/exon organization of the human 3-adrenergic receptor gene.--

5-175

Please replace paragraph [0022] with the following replacement paragraph:

--[0022] FIGURES 2A-B depict the nucleotide sequence (SEQ ID NO:3), amino acid translation (SEQ ID NO:4) and intron/exon organization of the mouse 3-adrenergic receptor gene.--

5-1143

Please replace paragraph [0025] with the following replacement paragraph:

--[0025] FIGURES 5A-B are a schematic representation of human and mouse adrenergic receptor mRNA splicing.--

5-14-57

Please replace paragraph [0032] with the following replacement paragraph:

--[0032] Besides polypeptides, the present invention also encompasses any nucleotide sequence of 23-adrenergic receptors in mammals. A preferred embodiment of these nucleotide sequences are encompassed in FIGS. 1A-B (SEQ ID NO:1) and 2A-B (SEQ ID NO:3). Variants of the nucleotide sequence are also encompassed in the present invention including mutations and point substitutions using the above-described mutagenesis methods, provided that these variations do not significantly alter 23-adrenergic receptor activity.--

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Please replace paragraph [0042] with the following replacement paragraph:

--[0042] More particularly, it is advantageous to use a full length probe having the nucleotide sequence as defined in FIGS. 1A-B (SEQ ID NO:1) and 2A-B (SEQ ID NO:3) to probe a genomic or cDNA library of different mammalian species to obtain the related 3-adrenergic receptor of interest. A fragment of the nucleotide probe can also be generated.--

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